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PATENT COOPERATION TREA

From the INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

To:

BURNSIDE, Man Eli Lilly And Company P. O. Box 6288

Indianapolis, IN 46206-6288 ETATS-UNIS D'AMERIQUE SEP 13 2004

ELI LILLY AND COMPANY
Patent Division

NOTIFICATION OF TRANSMITTAL OF THE INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Rule 71.1)

Date of mailing

(day/month/year)

06.09.2004

Applicant's or agent's file reference

X-15906 🗸

IMPORTANT NOTIFICATION

International application No. PCT/US 03/26304

International filing date (day/month/year)

Priority date (day/month/year)

22.09.2003

25.09.2002

Applicant

ELI LILLY AND COMPANY et al.

- The applicant is hereby notified that this International Preliminary Examining Authority transmits herewith the international preliminary examination report and its annexes, if any, established on the international application.
- A copy of the report and its annexes, if any, is being transmitted to the International Bureau for communication to all the elected Offices.
- 3. Where required by any of the elected Offices, the International Bureau will prepare an English translation of the report (but not of any annexes) and will transmit such translation to those Offices.

#### 4. REMINDER

The applicant must enter the national phase before each elected Office by performing certain acts (filing translations and paying national fees) within 30 months from the priority date (or later in some Offices) (Article 39(1)) (see also the reminder sent by the International Bureau with Form PCT/IB/301).

Where a translation of the international application must be furnished to an elected Office, that translation must contain a translation of any annexes to the international preliminary examination report. It is the applicant's responsibility to prepare and furnish such translation directly to each elected Office concerned.

For further details on the applicable time limits and requirements of the elected Offices, see Volume II of the PCT Applicant's Guide.

The applicant's attention is drawn to Article 33(5), which provides that the criteria of novelty, inventive step and industrial applicability described in Article 33(2) to (4) merely serve the purposes of international preliminary examination and that "any Contracting State may apply additional or different criteria for the purposes of deciding whether, in that State, the claimed inventions is patentable or not" (see also Article 27(5)). Such additional criteria may relate, for example, to exemptions from patentability, requirements for enabling disclosure, clarity and support for the claims.

Name and mailing address of the international preliminary examining authority:



European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d

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**Authorized Officer** 

Ullrich, J

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## **PCT**

### INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference X-15906	FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/PEA/416)				
International application No. PCT/US 03/26304	International filing date (day/mor 22.09.2003	Priority date (day/month/year) 25.09.2002				
International Patent Classification (IPC) or both national classification and IPC C07D409/12						
Applicant ELI LILLY AND COMPANY et al.						
<ol> <li>This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.</li> </ol>						
2. This REPORT consists of a total	of 5 sheets, including this cove	r sheet.				
been amended and are the	This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).					
These annexes consist of a total	These annexes consist of a total of 2 sheets.					
This report contains indications r	elating to the following items:					
I ⊠ Basis of the opinion						
II 🗆 Priority						
III 🖾 Non-establishment of	f opinion with regard to novelty, i	nventive step and industrial applicability				
IV 🔲 Lack of unity of inven	tion					
	V 🖾 Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement					
VI □ Certain documents ci	ited					
_	international application					
VIII   Certain observations on the international application						
Date of submission of the demand	Date o	completion of this report				
15.03.2004	06.09	.2004				
Name and mailing address of the internatio	nal Author	zed Officer				
preliminary examining authority:  European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523	656 epmu d	am Bakhtiary, M one No. +49 89 2399-8556				

## INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No.

PCT/US 03/26304

I. B	asis	of	the	re	po	rt
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1. With regard to the **elements** of the international application (Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17)):

	Des	Description, Pages					
	1-4	2	as originally filed				
	<b>0</b> 1.	· · · · · · · · · · · · · · · · · · ·					
		ims, Numbers					
		1, 22 (part)	as originally filed				
	22 (	(part), 23-39	received on 16.07.2004 with letter of 16.07.2004				
2.	With lang	Vith regard to the <b>language</b> , all the elements marked above were available or furnished to this Authority in thanguage in which the international application was filed, unless otherwise indicated under this item.					
	The	ese elements were av	ailable or furnished to this Authority in the following language: , which is:				
		the language of a tra	anslation furnished for the purposes of the international search (under Rule 23.1(b)).				
☐ the language of publication of the internation			lication of the international application (under Rule 48.3(b)).				
		the language of a tra Rule 55.2 and/or 55.	anslation furnished for the purposes of international preliminary examination (under 3).				
3.	Witl inte	With regard to any <b>nucleotide and/or amino acid sequence</b> disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:					
	☐ contained in the international application in written form.						
		filed together with the international application in computer readable form.					
		furnished subsequently to this Authority in written form.					
		furnished subsequer	ntly to this Authority in computer readable form.				
		The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.					
		The statement that the listing has been furn	he information recorded in computer readable form is identical to the written sequence ished.				
4.	. The amendments have resulted in the cancellation of:						
		the description,	pages:				
		the claims,	Nos.:				
		the drawings,	sheets:				
5.		☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)).					
		(Any replacement sh report.)	neet containing such amendments must be referred to under item 1 and annexed to this				
<b>3</b> .	Additional observations, if necessary:						

# INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No.

PCT/US 03/26304

111.	. No	n-establishment of opinion w	vith re	gard to nov	elty, inventive step and industrial applicability	
1.	The obv	ne questions whether the claimed invention appears to be novel, to involve an inventive step (to be non- vious), or to be industrially applicable have not been examined in respect of:				
☐ the entire international application,						
	⊠	claims Nos. 17-28				
		because:				
	×	the said international application, or the said claims Nos. 17-28 regarding industrial applicability relate to the following subject matter which does not require an international preliminary examination (specify):				
		see separate sheet				
	the description, claims or drawings (indicate particular elements below) or said claims Nos. are so unclear that no meaningful opinion could be formed (specify):					
the claims, or said claims Nos. are so inadequately supported by the description that no meaning could be formed.				ely supported by the description that no meaningful opinion		
		no international search report has been established for the said claims Nos.				
2. A meaningful international preliminary examination cannot be carried out due to the failure of the nucleor amino acid sequence listing to comply with the standard provided for in Annex C of the Administrativ Instructions:						
		the written form has not been furnished or does not comply with the Standard.				
☐ the computer readable form has not been furnished or does not comply with the Standard.				hed or does not comply with the Standard.		
V.		soned statement under Artic tions and explanations supp			ard to novelty, inventive step or industrial applicability; ment	
1.	Sta	tement				
	Nov	velty (N)	Yes: No:	Claims Claims	1-39	
	Inve	entive step (IS)	Yes: No:	Claims Claims	1-39	
	Indi	ustrial applicability (IA)	Yes: No:	Claims Claims	1-16,29-39 17-28	

2. Citations and explanations

see separate sheet

#### **EXAMINATION REPORT - SEPARATE SHEET**

#### Re Item III

Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

Claims 17-28 relate to subject-matter considered by this Authority to be covered by the provisions of Rule 67.1(iv) PCT. Consequently, no opinion will be formulated with respect to the industrial applicability of the subject-matter of these claims (Article 34(4)(a)(I) PCT).

#### Re Item V

Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

- 1. Reference is made to the following documents:
  - D1: GRESE T A ET AL: 'STRUCTURE-ACTIVITY RELATIONSHIPS OF SELECTIVE ESTROGEN RECEPTOR MODULATORS: MODIFICATIONS TO THE 2-ARYLBENZOTHIOPHENE CORE OF RALOXIFENE' JOURNAL OF MEDICINAL CHEMISTRY, AMERICAN CHEMICAL SOCIETY. WASHINGTON, US, vol. 40, no. 2, 17 January 1997 (1997-01-17), pages 146-167, XP002050782 ISSN: 0022-2623
  - D2: EP-A-0 759 434 (LILLY CO ELI) 26 February 1997 (1997-02-26)
  - D3: EP-A-0 761 669 (LILLY CO ELI) 12 March 1997 (1997-03-12)
  - D4: WO 02 16316 A (MESTRES JORDI ;AKZO NOBEL NV (NL); LOOZEN HUBERT JANJOZEF (NL); VE) 28 February 2002 (2002-02-28)
- 2. Novelty and inventive step (articles 33.2,33.3 PCT)

Regarding main claim 1, the closest prior art may be considered as being WO0216316 (D4).

The problem to be solved may be regarded as being to provide novel compounds useful as estrogen receptor modulators.

In view of the drastic structural differences (fused rings) the skilled person would not have predicted that such changes would leave the modified compound as biologically active.

Novelty and inventivity may be recognised for final compounds.

Claims related to intermediates have been deleted, therefore novelty objections made in our written opinion are not valid anymore.

# INTERNATIONAL PRELIMINARY International application No. PCT/US 03/26304 EXAMINATION REPORT - SEPARATE SHEET

### 3. Industrial application (article 33.4 PCT)

For the assessment of the present claims 17-28 on the question whether they are industrially applicable, no unified criteria exist in the PCT Contracting States. The patentability can also be dependent upon the formulation of the claims. The EPO, for example, does not recognize as industrially applicable the subject-matter of claims to the use of a compound in medical treatment, but may allow, however, claims to a known compound for first use in medical treatment and the use of such a compound for the manufacture of a medicament for a new medical treatment.

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